## What is claimed is:

- 1. A composition for the inhibition of the translation of a Mect1-MAML2 chimeric gene consisting essentially of: (a) a fragment of the nucleic acid encoding the Mect1-MAML2 chimeric gene, and (b) a nucleic acid complementary to the fragment, wherein the fragment is about 17 to about 32 nucleotides in length.
- 2. The composition of claim 1, wherein the nucleic acid molecule complementary to the fragment has from 1 to about 10 base substitutions, and wherein the nucleic acid molecule complementary to the fragment optionally contains one or more insertions and is about 19 to about 35 nucleotides in length.
- 3. The composition of claim 2, further consisting essentially of a restriction enzyme sequence, wherein the fragment and the nucleic acid complementary to the fragment are joined by the restriction enzyme sequence.
- 4. The composition of claim 1, wherein the Mect1-MAML2 chimeric gene has the nucleotide sequence of SEQ ID NO: 1.
- 5. The composition of claim 1, wherein the Mect1-MAML2 chimeric gene results from a t(11;19) translocation.
- 6. The composition of claim 2, wherein the fragment has the nucleotide sequence of SEQ ID NO: 5 or 6.
- 7. The composition of claim 6, wherein the nucleic acid complementary to the fragment has the nucleotide sequence of SEQ ID NO: 7.
  - 8. The composition of claim 1, wherein the composition is in a vector.

- 9. The composition of claim 8, wherein the vector is a plasmid.
- 10. The composition of claim 8, wherein the vector is a viral vector.
- 11. The composition of claim 10, wherein the viral vector is an adenoviral vector.
- 12. The composition of claim 3, wherein the fragment is about 21 to about 32 nucleotides in length.
- 13. The composition of claim 12, wherein the fragment is about 28 to about 29 nucleotides in length.
- 14. The composition of claim 3, wherein the restriction enzyme sequence is a *Hin* dIII sequence.
- 15. The composition of claim 2, wherein the nucleic acid molecule complementary to the fragment has from about 2 to about 5 substitutions.
- 16. The composition of claim 1, wherein the composition has the nucleotide sequence of SEQ ID NO: 2, 3, or 4.
- 17. The composition of claim 1, wherein the fragment has the nucleotide sequence of SEQ ID NO: 8 or 9.
- 18. The composition of claim 1, wherein the fragment is about 17 to about 22 nucleotides in length.
- 19. The composition of claim 18, wherein the fragment is about 19 to about 21 nucleotides in length.

- 20. The composition of claim 1, wherein the fragment and the nucleic acid complementary to the fragment are under the control of different promoters on the same nucleic acid molecule.
- 21. The composition of claim 20, wherein the promoters are RNA polymerase promoters.
- 22. The composition of claim 21, wherein the promoters are RNA polymerase III promoters.
- 23. The composition of claim 1, wherein, upon annealing of the transcripts of the fragment and the nucleic acid complementary to the fragment, the annealed transcripts of the composition have a 3' overhang consisting of 1 to about 4 nucleotides on one or both ends of the annealed transcripts.
- 24. The composition of claim 23, wherein the 3' overhang consists of about 2 to about 3 nucleotides.
- 25. The composition of claim 23, wherein one or more of the nucleotides of the 3' overhang are uridine.
- 26. The composition of claim 23, wherein the 3' overhang consists of 2 uridine residues.
- 27. A method of inhibiting the translation of a Mect1-MAML2 chimeric gene in a cell comprising contacting the cell expressing the Mect1-MAML2 chimeric gene with the composition of claim 1, whereupon the translation of the Mect1-MAML2 chimeric gene in the cell is inhibited.

- 28. The method of claim 27, wherein the cell comprises a t(11;19) translocation, wherein the translocation results in a Mect1-MAML2 chimeric gene.
  - 29. The method of claim 27, wherein the cell is in a host.
  - 30. The method of claim 29, wherein the host is a mammal.
  - 31. The method of claim 30, wherein the mammal is a human.
- 32. The method of claim 30, wherein the cell is a cancerous cell of mucepidermoid origin and the inhibition of the translation of the Mect1-MAML2 chimeric gene results in the inhibition of the cancerous cell.
  - 33. The method of claim 32, wherein the cancerous cell is in a gland.
  - 34. The method of claim 33, wherein the gland is a salivary gland.